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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/587,995	07/27/2007	Susan Marie Metcalfe	2655.0010000/RWE	7016
STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C. 1100 NEW YORK AVENUE, N.W.			EXAMINER	
			LOCKARD, JON MCCLELLAND	
WASHINGTON, DC 20005			ART UNIT	PAPER NUMBER
			1647	
			MAIL DATE	DELIVERY MODE
			09/17/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)	
	10/587,995	METCALFE, SUSAN MARIE	
Office Action Summary	Examiner	Art Unit	
	JON M. LOCKARD	1647	
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with the c	orrespondence address	
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING D. - Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period of Failure to reply within the set or extended period for reply will, by statute Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tin will apply and will expire SIX (6) MONTHS from a, cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).	
Status			
Responsive to communication(s) filed on 31 Ju This action is FINAL . 2b) ☐ This Since this application is in condition for alloward closed in accordance with the practice under E	s action is non-final. nce except for formal matters, pro		
Disposition of Claims			
4) ☐ Claim(s) 21-33 is/are pending in the applicatio 4a) Of the above claim(s) is/are withdray 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) 21-33 are subject to restriction and/or	wn from consideration.		
Application Papers			
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) acc Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Example 11.	epted or b) objected to by the I drawing(s) be held in abeyance. See tion is required if the drawing(s) is ob	e 37 CFR 1.85(a). lected to. See 37 CFR 1.121(d).	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Bureau * See the attached detailed Office action for a list	s have been received. s have been received in Applicati rity documents have been receive u (PCT Rule 17.2(a)).	on No ed in this National Stage	
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	ate	

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DETAILED ACTION

Election/Restrictions

1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 21-23, in so far as they are drawn to a method for inducing or regulating an immune response with an axotrophin polypeptide.

Group II, claim(s) 21-23, in so far as they are drawn to a method for inducing or regulating an immune response with an axotrophin polynucleotide.

Group III, claim(s) 23, in so far as it is drawn to a method for inducing or regulating an immune response with a substance that enhances the amount of an axotrophin polypeptide.

Group IV, claim(s) 23, in so far as it is drawn to a method for inducing or regulating an immune response with a substance that enhances the activity of an axotrophin polypeptide.

Group V, claim(s) 24, in so far as it is drawn to a method for inducing or regulating an immune response with a substance that decreases the amount of an axotrophin polypeptide.

Group VI, claim(s) 24, in so far as it is drawn to a method for inducing or regulating an immune response with a substance that decreases the activity of an axotrophin polypeptide.

Group VII, claim(s) 25 and 28-33, in so far as they are drawn to a method for determining the immune status of an individual comprising determining the level of an axotrophin polypeptide.

Group VIII, claim(s) 25 and 28-33, in so far as they are drawn to a method for determining the immune status of an individual comprising determining the level of an axotrophin polynucleotide.

Group IX, claim(s) 26 and 27, in so far as they are drawn to an isolated axotrophin polypeptide.

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Group X, claim(s) 26, in so far as it is drawn to an isolated axotrophin polynucleotide.

- 2. The inventions listed as Groups I-X do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Pursuant to 37 C.F.R. § 1.475(B-D), the ISA/US considers that where multiple products and processes are claimed, the main invention shall consist of the first invention of the category first mentioned in the claims and the first recited invention of each of the other categories related thereto. Accordingly, the main invention (Group I) comprises the first recited method, a method for inducing or regulating an immune response with an axotrophin polypeptide. Groups II-X do not share the same or corresponding special technical feature because the Group II invention is drawn to a method for inducing or regulating an immune response with an axotrophin polynucleotide, the Group III invention is drawn to a method for inducing or regulating an immune response with a substance that enhances the amount of an axotrophin polypeptide, the Group IV invention is drawn to a method for inducing or regulating an immune response with a substance that enhances the activity of an axotrophin polypeptide, the Group V invention is drawn to a method for inducing or regulating an immune response with a substance that decreases the amount of an axotrophin polypeptide, the Group VI invention is drawn to a method for inducing or regulating an immune response with a substance that decreases the activity of an axotrophin polypeptide, the Group VII invention is drawn to a method for determining the immune status of an individual comprising determining the level of an axotrophin polypeptide, the Group VIII invention is drawn to a method for determining the immune status of an individual comprising determining the level of an axotrophin polynucleotide, the Group IX invention is drawn to an isolated axotrophin polypeptide, and the Group X invention is drawn to an isolated axotrophin polynucleotide. Lack of unity is shown because these methods lack a common utility which is based upon a common technical feature which has been identified as the basis for that common utility.
- 3. Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement may be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.
- 4. The election of an invention or species may be made with or without traverse. To preserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

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invention to be rejoined.

- 5. The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and the product claims are subsequently found allowable, withdrawn process claims that depend from or otherwise require all the limitations of the allowable product claim will be considered for rejoinder. <u>All</u> claims directed to a nonelected process invention must require all the limitations of an allowable product claim for that process
- 6. In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103 and 112. Until all claims to the elected product are found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowable product claim will not be rejoined. See MPEP § 821.04(b). Additionally, in order to retain the right to rejoinder in accordance with the above policy, applicant is advised that the process claims should be amended during prosecution to require the limitations of the product claims. Failure to do so may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

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Advisory Information

Any inquiry concerning this communication or earlier communications from the examiner should be directed to **Jon M. Lockard**, **Ph.D.** whose telephone number is **(571) 272-2717**. The examiner can normally be reached on Monday through Friday, 8:00 AM to 4:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, **Manjunath N. Rao, Ph.D.**, can be reached on **(571) 272-0939**. The fax number for the organization where this application or proceeding is assigned is **571-273-8300**.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at **866-217-9197** (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Jon M. Lockard, Ph.D. September 15, 2008

/Jon M Lockard/ Examiner, Art Unit 1647